

Case Reports

evidence of the explanted valves, they concluded that 2 of the bioprostheses were thrombosed and that the other 2 had coagulated blood on the outflow of the cusps and thus should not be classified as demonstrating structural valve dysfunction. This view was not shared by Flameng and associates,⁴ however, who hypothesized that early stenotic-type valvular dysfunction in the 4 reported cases was perhaps related to patient–prosthesis mismatch or turbulent transvalvular flow. Their definition of moderate patient–prosthesis mismatch for patients with an indexed EOA of less than 0.85 has since been challenged by Jamieson and colleagues.⁵

We have presented the cases of 2 patients with early stenotic valve failure who did not have any evidence of patient–prosthesis mismatch, and had postoperative indexed EOAs of $0.90 \text{ cm}^2/\text{m}^2$ and $0.83 \text{ cm}^2/\text{m}^2$. In addition, the explanted valves did not reveal any pannus formation or histologic evidence of thrombosis, although thrombosis could not be ruled out in the first case on the basis of the available evidence. Importantly, there was no evidence of subtherapeutic anticoagulation or procoagulant states, and our

patients had normal renal function, without any documented hypercalcemic syndromes. After detailed pathologic examination, it remains unclear what caused the early valvular dysfunction of the Mosaic bioprostheses in our patients. In view of our findings, however, we believe that there is a need for clinical vigilance with regular echocardiographic follow-up in patients who have had implantation of Mosaic aortic bioprosthetic valves.

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Saphenous vein graft bronchopulmonary fistula after coronary artery bypass grafting presenting as chronic cough and subsequent massive hemoptysis

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Aortocoronary conduit bronchopulmonary fistula is a rare, potentially lethal complication seen after coronary artery bypass grafting (CABG). The ideal diagnostic modality in such cases is not established. We present a case of early-onset aortocoronary conduit bronchopulmonary fistula diagnosed by coronary angiography and immediate noncontrast computed tomography (CT) in a patient with massive hemoptysis 8 weeks after CABG.

CLINICAL SUMMARY

A 75-year-old woman underwent elective CABG. Conduits were fashioned from left internal thoracic artery to left anterior descending coronary artery and long saphenous vein graft to posterior descending coronary artery. The ascending aorta was noted to be thin walled and dilated, measuring 3.6 cm. On removal of the cardioplegia cannula from the ascending aorta, a complex 2-cm² tear was noted. Control of this area was undertaken by fashioning the proximal aorta–saphenous vein graft anastomosis in an oversized manner.

At follow-up, the patient was noted to have acquired a chronic cough. At 8 postoperative weeks, the patient had sudden significant hemoptysis. Contrast chest CT demonstrated a filling defect in the bronchus intermedius, consistent with hemorrhage, retrosternal fluid collection, and opacification within the medial portion of the right upper lobe (Figure 1, A). Rigid bronchoscopy showed large amounts of blood throughout the tracheobronchial tree but no active bleeding point. Coronary angiography

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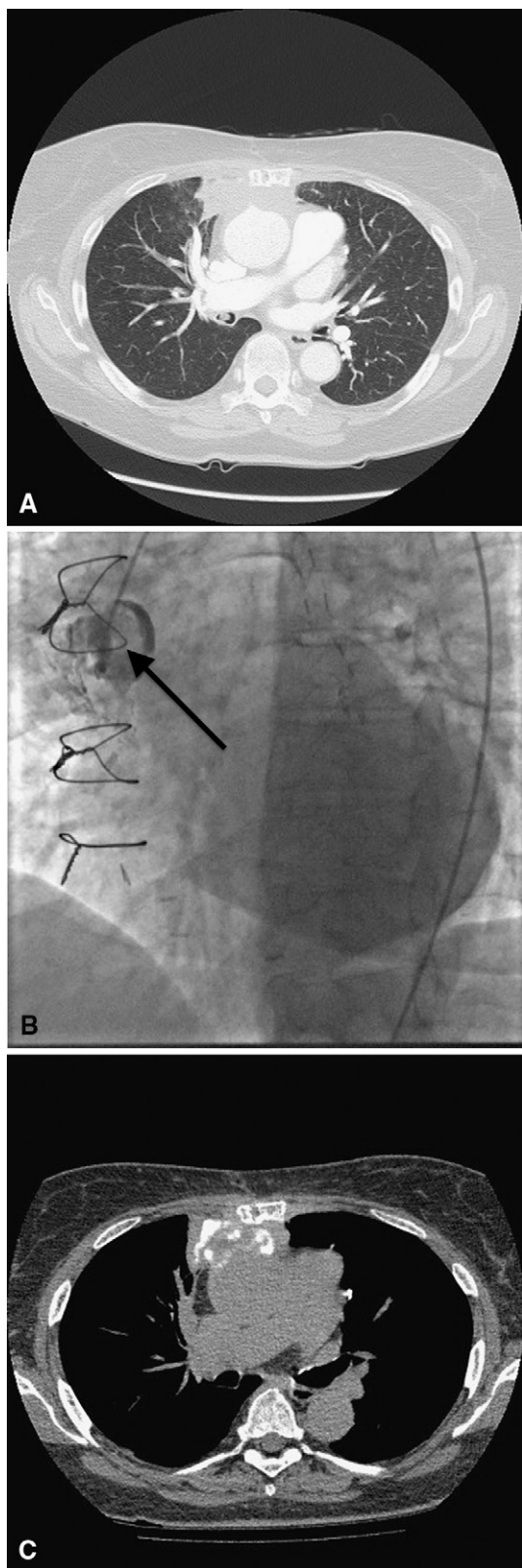


FIGURE 1. A, Contrast chest computed tomography demonstrating filling defect in the bronchus intermedius consistent with hemorrhage within airway, retrosternal fluid collection, and opacification of the right upper

lobe. B, Coronary angiography. Arrow demonstrates absence of flow within the saphenous vein graft to the right coronary artery. C, Noncontrast chest computed tomography performed immediately after coronary angiography demonstrating extravasation of contrast from the saphenous vein graft lying retrosternally and within the right upper lobe consistent with aortocoronary conduit fistula.

DISCUSSION

Aortocoronary conduit bronchopulmonary fistula is a rare but potentially lethal complication of CABG; however, the optimal diagnostic modality is unknown. We present a case of saphenous vein graft pulmonary fistula diagnosed by coronary angiography and immediate non-contrast chest CT in a patient with chronic cough and massive hemoptysis 8 weeks after CABG. Fistulous connections have been reported in association with both internal thoracic artery and saphenous vein grafts. Conduits may fistulate into the cardiac chambers,¹ pulmonary vasculature,² or the lung parenchyma.³ The most common presentation of aortocoronary conduit bronchopulmonary fistula is recurrent angina resulting from steal phenomenon; however, other reported modes of presentation include myocardial infarction resulting from distal embolization and congestive cardiac failure.⁴

Fistulas are usually although not exclusively associated with aneurysmal dilation of the graft and most commonly are seen 10 to 20 years after CABG¹ and in association with atretic grafts. In our case, symptomatic saphenous bronchopulmonary fistula was diagnosed only 8 weeks after CABG. This is to our knowledge the earliest presentation of such a fistula after CABG. Benchimol and colleagues⁵ proposed mechanisms of aneurysm formation inferred from the tendency of these lesions to occur in certain regions of the venous graft, including in the vicinity of

lobe. B, Coronary angiography. Arrow demonstrates absence of flow within the saphenous vein graft to the right coronary artery. C, Noncontrast chest computed tomography performed immediately after coronary angiography demonstrating extravasation of contrast from the saphenous vein graft lying retrosternally and within the right upper lobe consistent with aortocoronary conduit fistula.

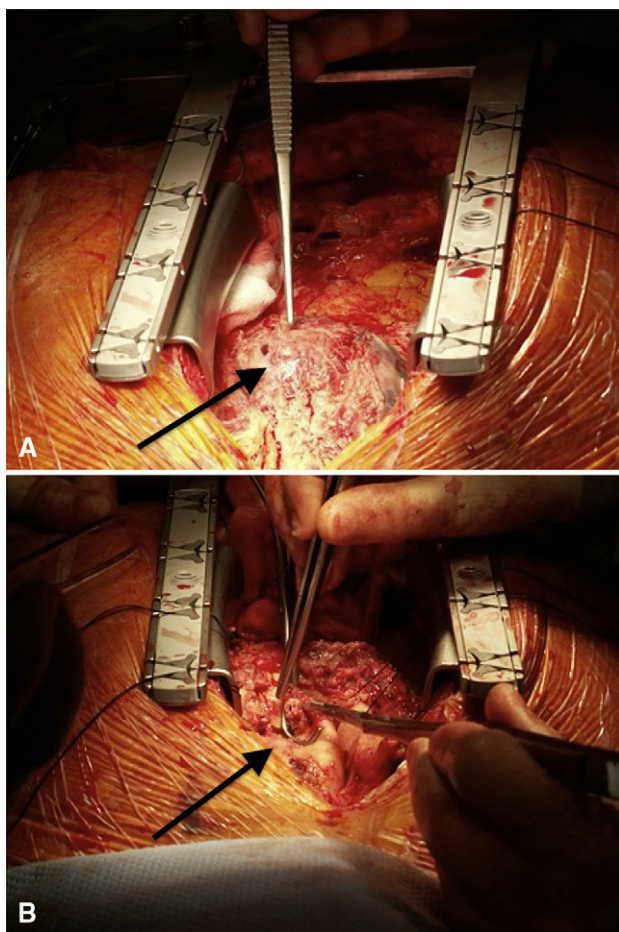


FIGURE 2. A, Intraoperative photograph. Arrow demonstrates retrosternal para-aortic collection at site of right coronary vein graft. B, Intraoperative photograph. Arrow shows fistula between saphenous vein graft (transected) and right lung.

valves because of the lack of circular muscle in these areas, at branch sites, in areas of iatrogenic injury, and at sites of atherosclerotic degeneration and vein wall necrosis or infection. In our case, it is likely that the etiology of fistula formation was related to the fashioning of an oversized

proximal anastomosis to deal with the aortic tear caused by the cardioplegia cannula, possibly in association with vein wall necrosis and infection as a result of application of surgical sealant.

Although in some cases fistulas may be asymptomatic, indications for treatment include refractory angina, heart failure, endocarditis, or, as in this case, hemoptysis. Surgical resection is the standard treatment; however, percutaneous coil embolization and covered stent deployment have also been successfully used.¹ Suggested diagnostic modalities include echocardiography, CT, magnetic resonance imaging, and angiography. Although bronchial angiography may be useful in cases of hemoptysis, it is limited to specialist centers, is technically challenging, and requires the patient to be actively bleeding at the time of angiography. In our case, we performed coronary angiography, which was not clearly diagnostic of a fistula. Angiography precipitated a further episode of hemoptysis, and subsequent noncontrast chest CT clearly showed contrast injected at coronary angiography present within the anterior mediastinum and airway of the right lung. We suggest that when aortocoronary graft fistula is suspected, the diagnostic algorithm of choice is coronary angiography immediately followed by noncontrast CT.

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